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(54) Title: TRANSTHORACIC DRUG DELIVERY DEVICE		
(57) Abstract		
<p>The present invention provides a transthoracic drug delivery device (10) that utilizes pressure to determine the precise location of the distal tip (18) of the delivery tube (14) of the device to insure that therapeutic substances are ejected into the myocardium (4) and not dissipated in unintended tissue locations. In one embodiment the drug delivery device (10) comprises a pressure sensing tube (12) mounted in parallel to a drug delivery tube (14) wherein the tubes are staggered so that the pressure tube (12) extends beyond the delivery tube (14). In another embodiment, a steerable syringe is provided so that multiple sites in the myocardium can be treated with a therapeutic substance with only one penetration into the heart. Once the delivery tube (28) of the syringe penetrates the left ventricle, its distal tip (32) can be deflected back toward the endocardial surface (6) and the tube (28) withdrawn proximally to cause penetration of the tip (32) into the myocardium (4). The deflectable tip (32) can be adjusted to a different deflection angle and the delivery tube (28) rotated to make available a circular array of points that may be accessed by the syringe.</p>		

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TRANSTHORACIC DRUG DELIVERY DEVICE

Field of the Invention

The present invention relates to a device and method for delivering a
5 therapeutic substance to the myocardium of a human heart.

Background of the Invention

Tissue becomes ischemic when it is deprived of adequate blood flow. Ischemia causes pain in the area of the affected tissue and, in the case of muscle
10 tissue, can interrupt muscular function. Left untreated, ischemic tissue can become infarcted and permanently non-functioning. Ischemia can be caused by a blockage in the vascular system that prohibits oxygenated blood from reaching the affected tissue area. However, ischemic tissue can be revived to function normally despite the deprivation of oxygenated blood because ischemic tissue can remain in a hibernating
15 state, preserving its viability for some time. Restoring blood flow to the ischemic region serves to revive the ischemic tissue.

Although ischemia can occur in various regions of the body, often tissue of the heart, the myocardium, is affected by ischemia due to coronary artery disease, occlusion of the coronary arteries, which otherwise provides blood to the myocardium. Muscle
20 tissue affected by ischemia can cause pain and lead to infarction of the tissue.

Ischemia can be treated, if a tissue has remained viable despite the deprivation of oxygenated blood, by restoring blood flow to the affected tissue.

Treatment of myocardial ischemia has been addressed by several techniques designed to restore blood supply to the affected region. Coronary artery bypass
25 grafting CABG involves grafting a venous segment between the aorta and the coronary artery to bypass the occluded portion of the artery. Once blood flow is redirected to the portion of the coronary artery beyond the occlusion, the supply of oxygenated blood is restored to the area of ischemic tissue.

Early researchers, more than thirty years ago, reported promising results for

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revascularizing the myocardium by piercing the muscle to create multiple channels for blood flow. Sen, P.K. et al., "Transmyocardial Acupuncture - A New Approach to Myocardial Revascularization", *Journal of Thoracic and Cardiovascular Surgery*, Vol. 50, No. 2, August 1965, pp. 181-189. Although others have reported varying degrees 5 of success with various methods of piercing the myocardium to restore blood flow to the muscle, many have faced common problems such as closure of the created channels. Various techniques of perforating the muscle tissue to avoid closure have been reported by researchers. These techniques include piercing with a solid sharp tip wire, hypodermic tube and physically stretching the channel after its formation.

10 Reportedly, many of these methods still produced trauma and tearing of the tissue that ultimately led to closure of the channel.

An alternative method of creating channels that potentially avoids the problem of closure involves the use of laser technology. Researchers have reported success in maintaining patent channels in the myocardium by forming the channels with the 15 heat energy of a laser. Mirhoseini, M. et al., "Revascularization of the Heart by Laser", *Journal of Microsurgery*, Vol. 2, No. 4, June 1981, pp. 253-260. The laser was said to form channels in the tissue that were clean and made without tearing and trauma, suggesting that scarring does not occur and the channels are less likely to experience the closure that results from healing. U.S. Patent No. 5,769843 (Abela et 20 al.) discloses creating laser-made TMR channels utilizing a catheter based system. Abela also discloses a magnetic navigation system to guide the catheter to the desired position within the heart. Aita patents 5,380,316 and 5,389,096 disclose another approach to a catheter based system for TMR.

Although there has been some published recognition of the desirability of 25 performing transmyocardial revascularization (TMR) in a non-laser catheterization procedure, there does not appear to be evidence that such procedures have been put into practice. For example, U.S. Patent No. 5,429,144 Wilk discloses inserting an expandable implant within a preformed channel created within the myocardium for the purposes of creating blood flow into the tissue from the left ventricle.

30 Performing TMR by placing stents in the myocardium is also disclosed in U.S.

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Patent No. 5,810,836 (Hussein et al.). The Hussein patent discloses several stent embodiments that are delivered through the epicardium of the heart, into the myocardium and positioned to be open to the left ventricle. The stents are intended to maintain an open channel in the myocardium through which blood enters from the 5 ventricle and perfuses into the myocardium.

Angiogenesis, the growth of new blood vessels in tissue, has been the subject of increased study in recent years. Such blood vessel growth provides new supplies of oxygenated blood to a region of tissue that has the potential to remedy a variety of tissue and muscular ailments, particularly ischemia. Primarily, the study has focused 10 on perfecting angiogenic factors such as human growth factors produced from genetic engineering techniques. It has been reported that injection of such a growth factor into myocardial tissue initiates angiogenesis at that site, which is exhibited by a new dense capillary network within the tissue. Schumacher et al., "Induction of Neo-Angiogenesis in Ischemic Myocardium by Human Growth Factors", *Circulation*, 1998; 15 97:645-650. The authors noted that such treatment could be an approach to management of diffused coronary heart disease after alternative methods of administration have been developed.

Mechanical devices have been disclosed for delivering therapeutic substances to the myocardium. International patent publication no. WO 98/05307 by Local Med, 20 Inc. discloses a method of using a conventional, rigid syringe to access the myocardium and left ventricle to deliver a therapeutic substance such as an angiogenic factor.

Summary of the Invention

25 The present invention provides a transthoracic drug delivery device that is specially configured to be precisely located in the myocardium for accurate placement of a therapeutic substance such as a drug. The device comprises a syringe having a delivery that is capable of penetrating the myocardium via the epicardium to access the left ventricle. Access to the heart is gained through the thorax. A pressure 30 monitor associated with the device indicates the position of the distal tip of the

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delivery tube. It is noted that, throughout the discussion of the invention in the specification, "distal" is meant to indicate the direction along the access path of the device leading internal to the patient and "proximal" indicates the direction along the access path leading external to the patient. Pressure may be monitored through the 5 delivery tube or through a separate pressure tube associated with the delivery tube. Specifically, the amount of pressure measured through the tube of the device informs the physician whether the distal port of the tube has penetrated the myocardium completely to reach the left ventricle. In the case of separate delivery and pressure tubes, the distance between the pressure tube opening and drug delivery tube 10 opening is known, so the user can determine whether the drug delivery tube opening is still within the myocardium, and, therefore, appropriately placed to inject a drug into the myocardium. Multiple drug delivery tubes may be associated with the device to perfuse the drug more quickly through multiple ports into the myocardium. It is noted that throughout the specification "drug" includes all varieties of therapeutic substances 15 that may be beneficial to the body, including pharmaceutical agents, genetically engineered substances or natural substances.

In another embodiment, a pressure sensing hypodermic tube, as in the first embodiment, is used to transthoracically access the heart and penetrate the left ventricle. However, the second embodiment is comprised of a single pressure 20 sensing tube. The distal portion of the pressure tube is flexible and made controllable or steerable by mechanisms well known in the art of catheter manufacture such as a pull wire bonded to the distal end of the tube and extending proximally for manipulation outside the patient. After penetrating the left ventricle, the single pressure tube may be bent into a "J" configuration and the shaft withdrawn proximally 25 to cause the curved tip of the J portion to penetrate the myocardium again, through the endocardial surface while moving in a proximal direction, similar to a fish hook. Entrance of the distal port of the pressure tube into the tissue results in a drop of measured pressure, indicating to the user that the distal tip of the pressure tube is within tissue, and, thus, appropriately placed to inject the drug or therapeutic 30 substance being carried by the device.

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After delivery of the therapeutic substance to a first location, the distal tip of the pressure tube may be backed out of the tissue by distal movement of the device into the ventricle and another location accessed. The steerable distal tip of the device permits the distal tip to be deflected through various angles relative to the longitudinal axis of the tube. Combining this range of movement with the rotational capability inherent in the device provides a locus of points about the penetration point into the ventricle where the distal tip of the needle can be placed into the myocardium to deliver therapeutic substances.

It is an object of the present invention to provide a drug delivery device that can access the myocardium and left ventricle of the heart transthoracically and accurately indicate the position of the distal tip of the device within the heart.

It is another object of the invention to provide a drug delivery device that is capable of delivering a therapeutic substance to the myocardium of the heart accurately and easily.

It is yet another object of the invention to provide a drug delivery device that is capable of delivering a therapeutic substance to the myocardium of the heart quickly via a plurality of delivery conduits.

It is yet another object of the invention to provide a drug delivery device that is capable of reaching a plurality of delivery locations in the myocardium with a single access to the heart.

It is yet another object of the invention to provide a drug delivery device for placement within the heart that is associated with a pressure sensor having adequate sensitivity to discern whether the pressure sensor port is within the myocardium or within the left ventricle.

It is yet another object of the invention to provide a delivery device that is configured to have a deflectable tip so that the reverse side of tissue having been accessed by the device can also be penetrated by the sharp tip of the device as the device is moved back along its access path.

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The foregoing and other objects and advantages of the invention will be appreciated more fully from the following further description thereof, with reference to the accompanying diagrammatic drawings wherein:

FIG. 1 is a diagrammatic, sectional illustration of the left ventricle of the human heart having the drug delivery device of the present invention inserted through the myocardial tissue;

FIG. 2 is a detailed sectional illustration of a portion of myocardial tissue that has been penetrated by the drug delivery device of the present invention;

FIG. 3 is a detailed sectional illustration of a portion of myocardial tissue that has been penetrated by an alternate embodiment of the drug delivery device of the present invention;

FIG. 4 is a diagrammatic, sectional illustration of the left ventricle of the human heart wherein the myocardium has been penetrated by a second embodiment of the drug delivery device; and

FIGS. 5-8 show a series of detailed sectional illustrations of a portion of the myocardium having been penetrated by the second embodiment of the delivery device.

Description of the Illustrative Embodiment

20

FIG. 1 shows a sectional view of the left ventricle of the heart having been penetrated by the drug delivery device of the present invention. The multiple tube embodiment 10 is comprised of a pressure tube 12 for monitoring pressure and a delivery tube 14 for dispensing the therapeutic substance or drug into the myocardium. The tubes may comprise metal hypodermic tubing or any tubing capable of being configured to have sufficient strength and sharpness to penetrate tissue. The tubes are joined together side-by-side in parallel relationship, but are offset so that the pressure tube extends beyond the delivery tube. Both the pressure tube 12 and delivery tube 14 have sharpened distal tips 16 and 18 and open distal ports 20 and 22, respectively. The device is delivered through the thorax and epicardial surface 8

of the heart and advanced through the myocardium 4 until the pressure tube 12 penetrates the left ventricle 2. Alternatively, as will be described in further detail below, the tubes may be formed from relatively flexible material having rigid sharp inserts at the distal ends and the flexible tubes being temporarily fortified to resist
5 buckling during delivery by an inner stiffening element that may be withdrawn after delivery.

The proximal end of the delivery tube 14 is joined to a pressurized source of a therapeutic substance such as may be provided by a syringe (not shown). The pressure tube is joined at its proximal end 24 to a pressure monitor (not shown) for
10 measuring the pressure sensed at the pressure tube distal port 20. As the pressure tube is inserted through the tissue of the heart, measured pressure will be low; however, once the pressure port 20 reaches the left ventricle 2, measured pressure increases dramatically signaling the physician that the ventricle has been reached.

The distance D between the relative longitudinal spacing of the pressure tube
15 and delivery tube (measured from distal tip 16 to distal tip 18) is known. Therefore, at the moment during advancement of the device into the heart when pressure dramatically increases, not only is the position of distal tip 16 known (having penetrated the endocardial surface 6 to access the ventricle 2) but also, the position of the delivery tube distal tip 18 will be known relative to the endocardial surface 6.
20 Therefore, if it is desired to deliver a therapeutic substance to a certain depth beneath the endocardial surface 6 of the myocardium 4, the pressure and delivery tubes 12 and 16 can be arranged relative to each other to provide that distance D between their distal tips 16 and 18. Upon penetration of the ventricle, the therapeutic substance can be discharged from the delivery port 22 at the desired distance
25 beneath the endocardial surface 6. During this procedure, the pressure tube 12 may also be used to inject contrast media into the ventricle to help visualize the position of the device. FIG. 3 shows a variation of the multiple tube embodiment employing a plurality of delivery tubes, each spaced a distance beneath the distal tip of the pressure tube 12. The purpose of multiple delivery tubes is to provide increased drug

delivery capability to surrounding myocardial tissue 4 with each penetration of the device into the heart.

The second embodiment of the drug delivery device further addresses the desirability of minimizing the number of times that the device must be inserted into the heart. A deflectable syringe tube 28 is shown in FIG. 4 protruding through the myocardium 4 into the left ventricle 2 of a human heart 1. After penetrating the myocardium, the distal portion 30 of the tube 28 may be deflected away from the longitudinal axis of the tube by as much as approximately 180°. The objective of providing such a large deflection is to permit the distal tip 32 of the tube to penetrate the endocardial surface 6 as the tube is withdrawn in a proximal direction from the ventricle 2. In the deflected configuration, the distal portion 30 of the tube is J-shaped and, therefore, pierces the tissue in similar fashion to a fishhook as it is withdrawn proximally. Once the distal port 34 has penetrated the myocardium, the therapeutic substance can be ejected through the port and into the tissue. The primary benefit provided by the deflectable tip tube is that multiple tissue locations can be reached with a single insertion penetration of the device into the heart.

As shown in FIG. 5, the tube should be inserted into the myocardium 4 while in a straight configuration. However, to be deflectable, at least the distal portion 30 of the tube must be flexible. Therefore, to provide sufficient longitudinal support during delivery through the heart tissue, a stiffening element 36 such as a stainless steel wire, may be advanced through the lumen 38 of the tube, extending to the distal tip 32. At least the proximal portion 30 of the tube should be fabricated from a relatively flexible material, such as a relatively low density polymer, to provide adequate flexibility for bending when the tip is deflected. The distal tip 32 may comprise an insert of a more rigid material bonded to the flexible material, such as a high density polymer or stainless steel to provide a sharp, piercing profile.

As shown in FIG. 6, after penetrating the myocardium 4, the stiffening element 36 is withdrawn proximally somewhat from the distal portion 30 of the tube 28 so that the tip 32 may be deflected to present its piercing distal tip 32 toward the endocardial surface 6 of the myocardium 4. The stiffening element 36 may, but need not be

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- withdrawn completely from the tube to deflect the tip. The tip deflection may be actuated by a pull wire extending through the tube 28 and joined to the area of the distal tip 32. The pull wire extends proximally, out of the device so that it may be manipulated by the physician external to the patient. Pulling the pull wire puts tension
5 on one side of the tube which tends to cause the flexible distal portion 30 to curve in the direction that the tension is being applied. Such pull wire arrangements are well known in the catheter art and are not discussed in detail here.

- As shown in FIG. 7, the deflected tip is drawn proximally into the endocardial surface 6. In this configuration, the therapeutic substance is ejected from the port 34 into the tissue 4. After the desired quantity of therapeutic substance has been delivered to the location, the distal tip 32 is withdrawn from the myocardium 4 by moving the tube 28 distally slightly. The lumen of the tube 34 is in communication with a pressure monitor as was described in connection with the previous embodiment. Pressure sensing is useful in the present embodiment so that the
10 physician can determine when the distal tip 32 of the tube has penetrated the myocardium 4 so that drugs can be delivered through port 34. In use, the pressure monitor indicates a high pressure when the port 32 is open to the ventricle. However, the pressure reading drops appreciably once the distal port 34 has become submerged within tissue indicating to the user that the distal tip is implanted.
15
20 As shown in FIG. 8, the withdrawn distal tip 32 and distal portion 30 may be adjusted to locate a new tissue site in the myocardium to deliver the therapeutic substance. The amount of deflection of the distal tip, represented by arrow 40 may be adjusted to reach a new location within the myocardium 4 into which a therapeutic substance needs to be delivered. Additionally, the tube 28 may be rotated, as
25 indicated by arrow 46 to provide a circle of points that are accessible by the deflectable tip tube. A possible new position for the distal portion 30 of the tube is shown in phantom in FIG. 8 to exhibit the steering capabilities of the device.
30 From the foregoing it will be appreciated that the invention provides a drug delivery device and method for delivering therapeutic substances to the myocardium quickly and accurately by accessing the thorax with a syringe tube device.

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Additionally, the present invention provides a device and method for accessing multiple points on the endocardial surface of the myocardium with a single penetration into the heart.

It should be understood, however, that the foregoing description of the
5 invention is intended merely to be illustrative thereof and that other modifications,
-- embodiments and equivalents may be apparent to those skilled in the art without
departing from its spirit. Having thus described the invention, what we desire to claim
and secure by Letters Patent is:

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CLAIMS

1. A transthoracic drug delivery device comprising:
a pressure sensing tube and a drug delivery tube joined to the pressure
5 sensing tube in a parallel relationship, but staggered so that one tube extends beyond
the other by a known distance.

2. A therapeutic drug delivery device as defined in claim 1 further
comprising:
10 a pressure sensor joined to the pressure sensing tube and a source of a
therapeutic substance joined to the drug delivery tube.

3. A therapeutic drug delivery device as defined in claim 1 wherein the
known distance between the ends of the staggered tubes is less than the thickness of
15 a subject patient's myocardium.

4. A transthoracic drug delivery device as defined in claim 1 wherein the
pressure sensing tube extends beyond the drug delivery tube.

- 20 5. A thoracic drug delivery device as defined in claim 1 further comprising
a plurality of drug delivery tubes.

6. A method of delivering a therapeutic substance to the myocardium
comprising:
25 providing a delivery device comprised of a pressure sensing tube joined
to at least one drug delivery tube maintained in parallel but staggered so that one
extends beyond the other by a known distance that is less than the thickness of the
myocardium;
Transthoracically accessing the heart;
30 inserting the tubes into the heart;

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monitoring the pressure sensed by the tube while moving the device further into the heart;

- stopping advancement of the device into the heart when a dramatic pressure increase is noted indicating pressure sensing tube has entered the ventricle
- 5 and the delivery tube is within the myocardium;
- ejecting a quantity of a therapeutic substance from the delivery tube into the myocardium.

7. A steerable syringe comprising:

- 10 a delivery tube having a sharpened distal tip, a flexible distal portion, a distal tip, a port and a pull wire for deflecting the distal tip.

8. A transthoracic drug delivery device as defined in claim 7 further comprising a stiffening element slidable within the tube to provide support for the
15 flexible distal portion during delivery of the tube into the heart.

9. A method of delivering a therapeutic substance to a plurality of sites in the myocardium with a single penetration into the heart comprising:

- introducing a tube having a deflectable distal portion and distal tip into
20 the left ventricle of the heart;
- deflecting the distal tip of the tube so that the tip faces the endocardium;
- moving the tube in a proximal direction so that the distal tip penetrates
the endocardium;
- ejecting a therapeutic substance through the tube into the myocardium;
- 25 moving the distal tip from the myocardium by advancing the tube distally;
- adjusting the position of the tip by changing the deflection angle of the tip or by rotating the tube to a new myocardial location;
- moving the tube in a proximal direction to insert the distal tip into the
30 myocardium at the new location;

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ejecting a therapeutic substance from the tube into the myocardium.

10. A method of delivering a therapeutic substance to a plurality of locations within a myocardium as defined in claim 9 wherein the tube is equipped with a
5 pressure sensor and the presence of the distal tip within myocardial tissue is determined by the change in pressure between the pressure indicated when the distal tip is in the myocardium versus the pressure indicated when the distal tip was in the ventricle.
- 10 11. A transthoracic drug delivery device comprising:
a pressure sensing tube and a drug delivery tube joined to the pressure sensing tube.

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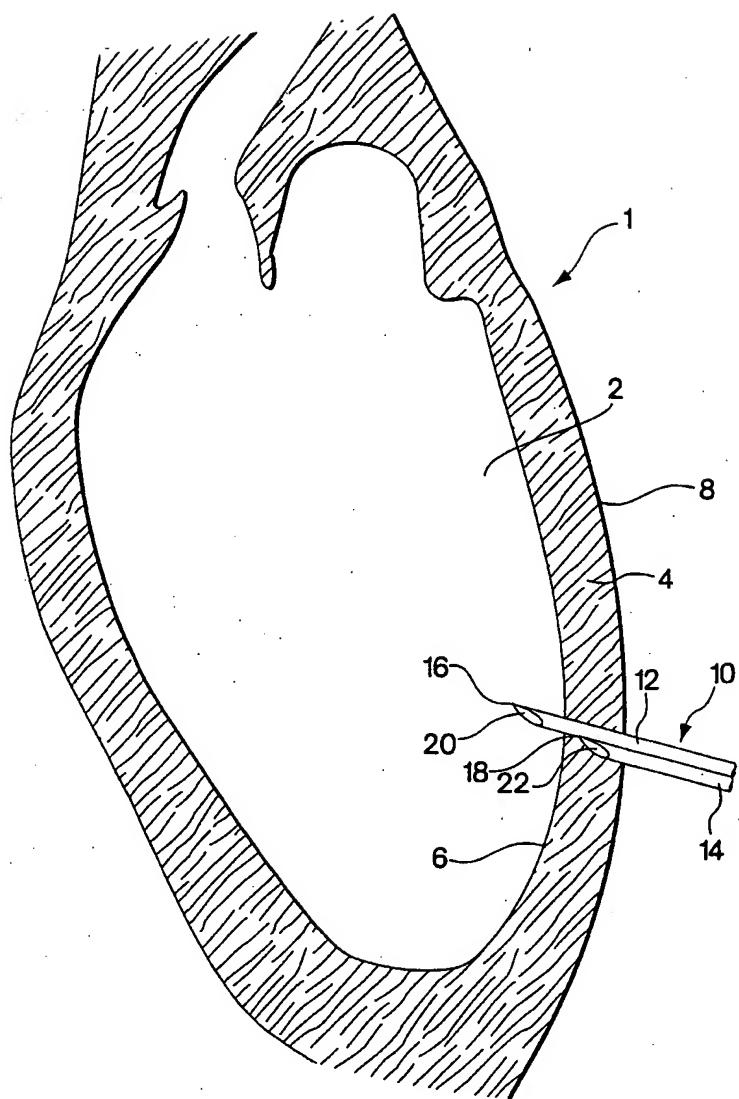


Fig. 1

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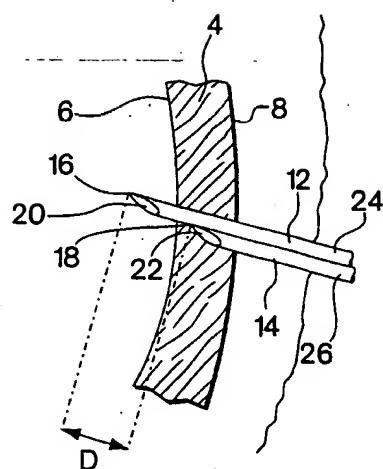


Fig. 2

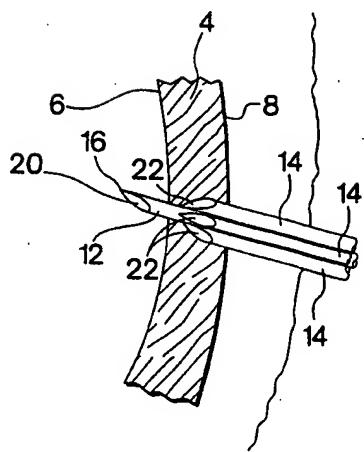


Fig. 3

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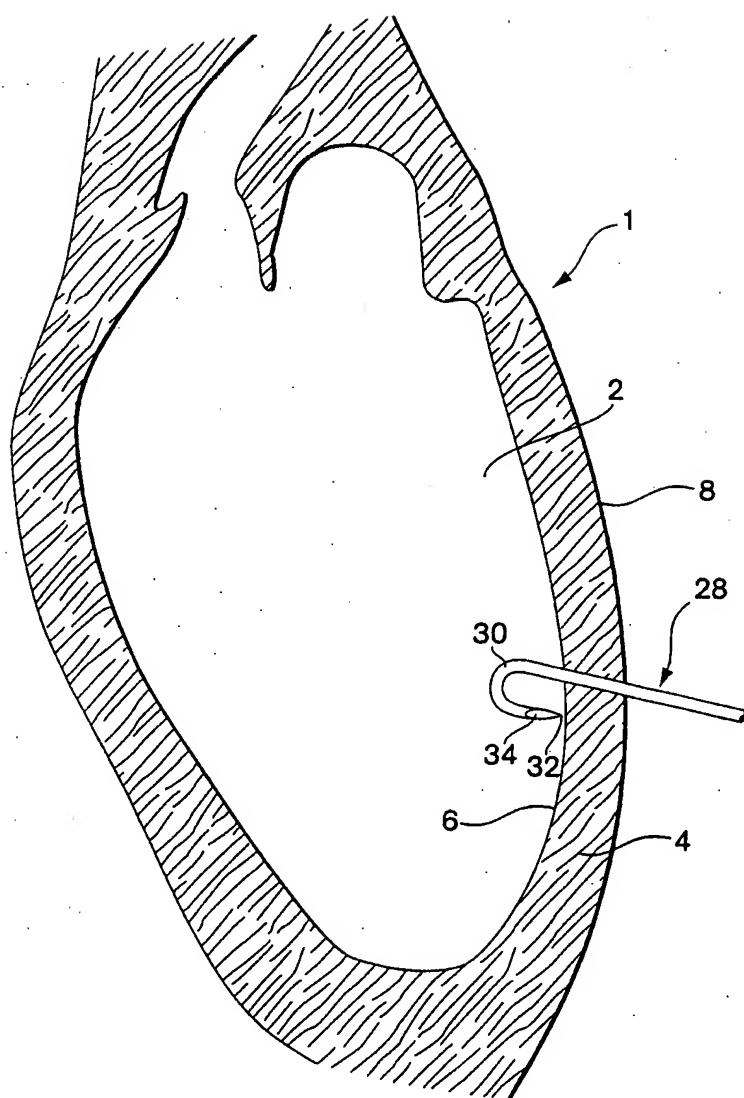


Fig. 4

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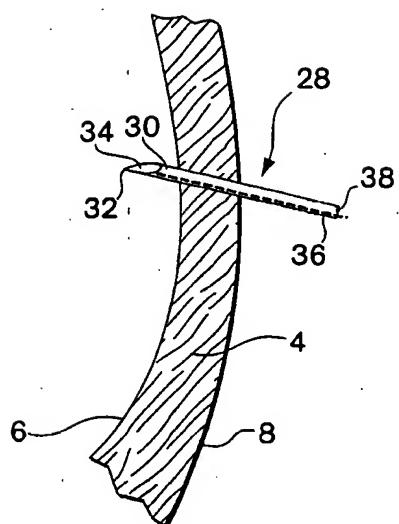


Fig. 5

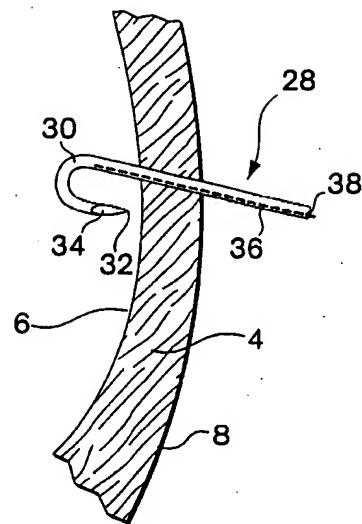


Fig. 6

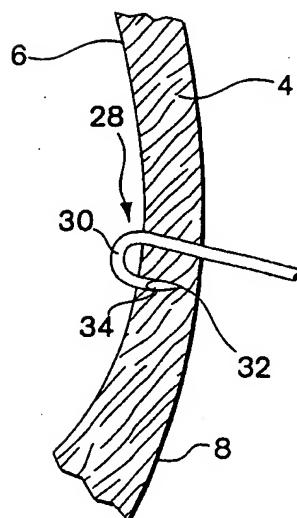


Fig. 7

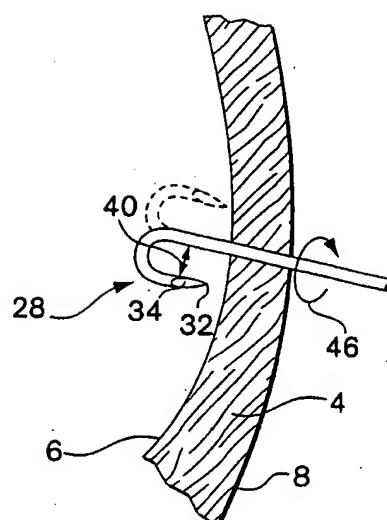


Fig. 8



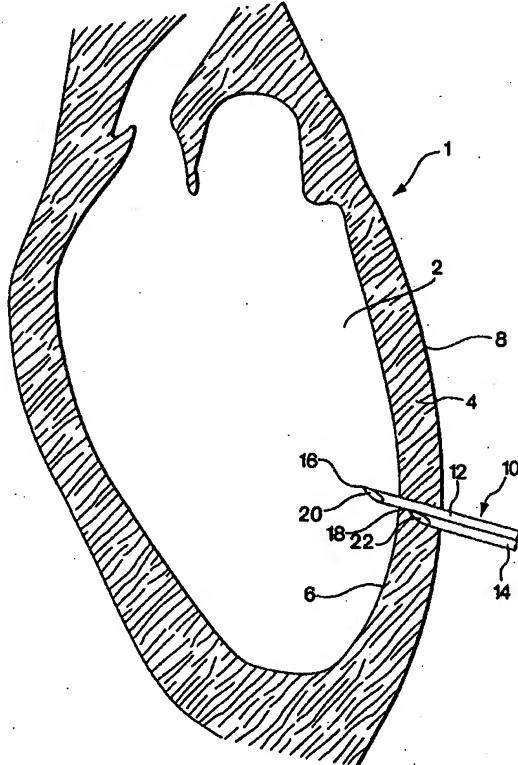
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(54) Title: TRANSTHORACIC DRUG DELIVERY DEVICE

(57) Abstract

The present invention provides a transthoracic drug delivery device (10) that utilizes pressure to determine the precise location of the distal tip (18) of the delivery tube (14) of the device to insure that therapeutic substances are ejected into the myocardium (4) and not dissipated in unintended tissue locations. In one embodiment the drug delivery device (10) comprises a pressure sensing tube (12) mounted in parallel to a drug delivery tube (14) wherein the tubes are staggered so that the pressure tube (12) extends beyond the delivery tube (14). In another embodiment, a steerable syringe is provided so that multiple sites in the myocardium can be treated with a therapeutic substance with only one penetration into the heart. Once the delivery tube (28) of the syringe penetrates the left ventricle, its distal tip (32) can be deflected back toward the endocardial surface (6) and the tube (28) withdrawn proximally to cause penetration of the tip (32) into the myocardium (4). The deflectable tip (32) can be adjusted to a different deflection angle and the delivery tube (28) rotated to make available a circular array of points that may be accessed by the syringe.



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CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

INTERNATIONAL SEARCH REPORT

International Application No PCT/US 99/22541

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 A61M25/00
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According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) IPC 7 A61M A61B
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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)
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C. DOCUMENTS CONSIDERED TO BE RELEVANT
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Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 490 459 A (KENDALL & CO) 17 June 1992 (1992-06-17) column 3, line 1 -column 5, line 42; figures 1-5 ---	1,2,4,5, 11
X	US 4 894 057 A (HOWES RANDOLPH M) 16 January 1990 (1990-01-16) column 3, line 25 -column 6, line 26; figures 1-7 ---	1,2,4,5, 11
A	column 3, line 25 -column 6, line 26; figures 1-7 ---	3
X	WO 90 06723 A (ENDOSONICS CORP) 28 June 1990 (1990-06-28) page 15, line 29 -page 18, line 33; figures 1-3C ---	1,2,4,11
	-/-	

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"Z" document member of the same patent family

Date of the actual completion of the international search

Date of mailing of the international search report
--

24 January 2000

03.05.2000

Name and mailing address of the ISA

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Authorized officer

Levert, C

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/22541

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 680 544 A (SHINNICK JAMES P ET AL) 1 August 1972 (1972-08-01) column 3, line 37 -column 6, line 28; figures 1-9 ---	1,11
A	US 5 429 144 A (WILK PETER J) 4 July 1995 (1995-07-04) cited in the application column 6, line 35 - line 66; figures 3A-4 -----	1,11

International application No.

PCT/US 99/22541

INTERNATIONAL SEARCH REPORT

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 6, 9, 10
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy
2. Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-5, 11

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US 99/22541

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-5, 11

A transthoracic drug delivery device comprising a drug delivery tube joined to a pressure sensing tube. (searched)

2. Claims: 7,8

A steerable syringe comprising a delivery tube having a sharpened distal tip, a flexible portion, a port and a pull wire for deflecting the distal tip. (not searched)

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/22541

Patent document cited in search report	Publication date	Patent family member(s)			Publication date
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US 3680544 A	01-08-1972	NONE			
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